

This is an official MS Health Alert Network (HAN) Alert

MESSAGE ID: MSHAN-20220303-00562-ALT (Health Alert)

RECIPIENTS: All Physicians, Hospitals, ERs, ICPs, NPs, PAs, and

Healthcare Providers – Statewide

Thursday, March 3, 2022

SUBJECT: UPDATE: Bebtelovimab Monoclonal Antibody for the

Treatment of Mild to Moderate COVID-19 Infection

UPDATES

• On Friday, February 11, 2022, the FDA announced Emergency Use Authorization (EUA) for Eli Lilly's bebtelovimab, a monoclonal antibody product now authorized for the treatment of mild to moderate COVID-19 in adults and certain pediatric patients who are at high risk for progression to severe COVID-19.

- The EUA for sotrovimab has also recently been updated to reduce the treatment window to administration of therapy within 7 days of symptom onset.
- Please see the following resources for utilization of bebtelovimab and other therapeutics for COVID-19 treatment:
 - NIH COVID-19 Treatment Guidelines: <u>Therapies | COVID-19 Treatment Guidelines (nih.gov)</u>
 - Role of bebtelovimab for the treatment of high-risk, non-hospitalized patients with mild to moderate COVID-19: <u>Statement on Bebtelovimab</u> | COVID-19 Treatment Guidelines (nih.gov)
 - o Therapeutics Decision Aid (phe.gov)

Bebtelovimab and sotrovimab are monoclonal antibody (mAb) medications authorized by the Food and Drug Administration (FDA) under an Emergency Use Authorization (EUA) for the treatment of COVID-19. The mAbs are prescribed by authorized healthcare providers based on patient risk assessment. The United States Government has purchased these medications and allocates them on a weekly basis to states and jurisdictions.

To request supply of bebtelovimab and/or sotrovimab, please submit the MSDH Weekly Monoclonal Antibody Data Request survey on Tuesdays by 2:00 PM. If you need access to this survey, email C19Therapeutics@msdh.ms.gov.

Healthcare Provider fact sheets:

- Sotrovimab Fact Sheet for HCP (fda.gov)
- Bebtelovimab Fact Sheet for HCP (fda.gov)



Patients receiving monoclonal antibody therapy must be given a copy of the "Fact Sheet for Patients, Parents, and Caregivers" for the product they are receiving:

- GSK Sotrovimab Fact Sheet for Patients, Parents, and Caregivers 02232022 (fda.gov)
- Fact Sheet for Patients, Parents and Caregivers Emergency Use Authorization (EUA) of Bebtelovimab for Coronavirus Disease 2019 (COVID-19) (fda.gov)

Please email <u>C19Therapeutics@msdh.ms.gov</u> with any questions or concerns.



Alerting Message Specification Settings

Originating Agency: Mississippi State Department of Health
Alerting Program: MS Health Alert Network (MS HAN)
Message Identifier: MSHAN-20220303-00562-ALT

Program (HAN) Type: Health Alert
Status (Type): Actual ()
Message Type: Alert

Reference: MSHAN-00562

Severity: Unknown

Acknowledgement: No

Sensitive: Not Sensitive
Message Expiration: Undetermined
Urgency: Undetermined
Delivery Time: 600 minutes

Definition of Alerting Vocabulary and Message Specification Settings

Originating Agency: A unique identifier for the agency originating the alert.

Alerting Program: The program sending the alert or engaging in alerts and

communications using PHIN Communication and Alerting (PCA)

as a vehicle for their delivery.

Message Identifier: A unique alert identifier that is generated upon alert activation

(MSHAN-yyymmdd-hhmm-TTT (ALT=Health Alert,

ADV=Health Advisory, UPD=Health Update,

MSG/INFO=Message/Info Service).

Program (HAN) Type: Categories of Health Alert Messages.

Health Alert: Conveys the highest level of importance; warrants immediate

action or attention.

Health Advisory: Provides important information for a specific incident or situation;

may not require immediate action.

Health Update: Provides updated information regarding an incident or situation;

unlikely to require immediate action.

Health Info Service: Provides Message / Notification of general public health

information; unlikely to require immediate action.

Status (Type):

Actual: Communication or alert refers to a live event Exercise: Designated recipients must respond to the

communication or alert

Test: Communication or alert is related to a technical,

system test and should be disregarded

Message Type:

Alert: Indicates an original Alert

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Update: Indicates prior alert has been Updated and/or superseded

Cancel: Indicates prior alert has been cancelled Error: Indicates prior alert has been retracted

Reference: For a communication or alert with a Message Type of "Update" or "Cancel", this attribute contains the unique Message Identifier of the original communication or alert being updated or cancelled. "n/a" = Not Applicable.

Severity:

Extreme: Extraordinary threat to life or property
Severe: Significant threat to life or property
Moderate: Possible threat to life or property
Minor: Minimal threat to life or property
Unknown: Unknown threat to life or property

Acknowledgement: Indicates whether an acknowledgement on the part of the recipient is required to confirm that the alert was received, and the timeframe in which a response is required (Yes or No).

Sensitive:

Sensitive: Indicates the alert contains sensitive content

Not Sensitive: Indicates non-sensitive content

Message Expiration: Undetermined.

Urgency: Undetermined. Responsive action should be taken immediately.

Delivery Time: Indicates the timeframe for delivery of the alert (15, 60, 1440,

4320 minutes (.25, 1, 24, 72 hours)).